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1,223 Deaths Reported After Pfizer Vaccine in First 3 Months of Rollout, Documents Reveal

A document released by the U.S. Food and Drug Administration shows Pfizer's own analysis of adverse events following its COVID vaccine revealed safety concerns yet the FDA refuses to acknowledge them.

By [Madhava Setty, M.D.](#)

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EDITOR'S NOTE (March 11, 2022): The most [recent release](#) of Pfizer COVID-19 vaccine documents sparked a flurry of news stories. Many of them referenced data released previously (November 2021) showing Pfizer reported to the U.S. Food and Drug Administration that the drugmaker was aware of 1,223 reports of deaths following the vaccine. The deaths, a subset of 42,086 case reports listing 158,893 adverse events, occurred between Dec. 1, 2020, and Feb. 28, 2021. The Defender first reported on these numbers and other revelations from the documents released in November in the article below.

In August, Public Health and Medical Professionals for Transparency (PHMPT) submitted a Freedom of Information Act (FOIA) request to the U.S. Food and Drug Administration (FDA) for all of the data within Pfizer's [COVID-19](#) vaccine biological product file — a body of information comprising some 329,000 pages.

The FDA, [arguing](#) its poorly staffed Center for Biologics Evaluation and Research did not have the capacity to quickly redact legally exempt material, such as Pfizer proprietary information and personal private information of trial participants, the agency asked to be allowed to release only 500 pages of this data per month, thus necessitating [55 years](#) for full disclosure.

The agency later requested up to [75 years](#) to complete the task. As of Nov. 17, only a fraction of the data in question had been released.

Here I will discuss one of these released documents, the "[Cumulative Analysis of Post-authorization Adverse Event Reports](#)." This document constitutes one part of Pfizer's responsibility for pharmacovigilance with respect to their Biological License Agreement with the FDA.

Pharmacovigilance refers to the science and activities relating to the detection, assessment, understanding and prevention of [adverse effects](#) or any other medicine-related problem.

Before we examine the quantity, seriousness and nature of the adverse events included in this document it is worthwhile to pause and consider just how significant this report should have been to the public.

BUY TODAY: Robert F. Kennedy, Jr.'s New Book — 'The Real Anthony Fauci'

Pfizer's vaccine had yet to complete full safety and efficacy testing, yet its product was being rapidly deployed on a healthy population that dwarfed the size of the vaccine's clinical trial.

The FDA and Pfizer were well aware that very real risks, if they existed, could not have been identified from the trials alone. There weren't enough participants, and the participants had not been observed for very long.

Everything may seem okay if you experiment on 20,000 people, but what happens when you experiment on a million people?

The "Cumulative Analysis of Post-authorization Adverse Event Reports" should have been the "everything looks good so far" reassurance the FDA was seeking. Why was it necessary to impel the FDA to make this information public through a court order?

In the discussion section of the document (section 4), Pfizer assures the FDA it "... performs frequent and rigorous signal detection on BNT162b2 cases."

What does "rigorous" signal detection mean? Did Pfizer survey a large number of vaccine recipients for adverse events and investigate them? No, it didn't.

This report is merely a compilation of unsolicited, in other words, passive, reports of adverse events directly brought to Pfizer's attention by recipients, cases reported by the health authorities, cases published in the medical literature, cases from Pfizer-sponsored marketing programs, non-interventional studies and cases of serious adverse events reported from clinical studies regardless of causality assessment.

In the report, Pfizer admitted the "magnitude of underreporting is unknown."

It is well accepted that passive reporting will inescapably lead to underreporting. Nevertheless, according to Pfizer's report:

"Due to the large numbers of spontaneous adverse event reports received for the product, the MAH (Marketing Authorisation Holder) has prioritised the processing of serious cases, in order to meet expedited regulatory reporting timelines and ensure these reports are available for signal detection and evaluation activity."

The authors continued:

"Pfizer also taken a [sic] multiple actions to help alleviate the large increase of adverse event reports. This includes significant technology enhancements, [sic] and process and workflow solutions, as well as increasing the number of data entry and case processing colleagues."

In other words, the number of adverse events reported overwhelmed Pfizer's expectations, yet the vaccine maker concluded, "The findings of these signal detection analyses are consistent with the known safety profile of the vaccine."

This paradoxical statement will prove to be an important clue as we dissect the data below.

What does the document reveal?

Through Feb. 28, a total of 42,086 recipients (cases) reported 158,893 events, or adverse reactions to the Pfizer vaccine. Approximately 50% of these events were deemed serious.

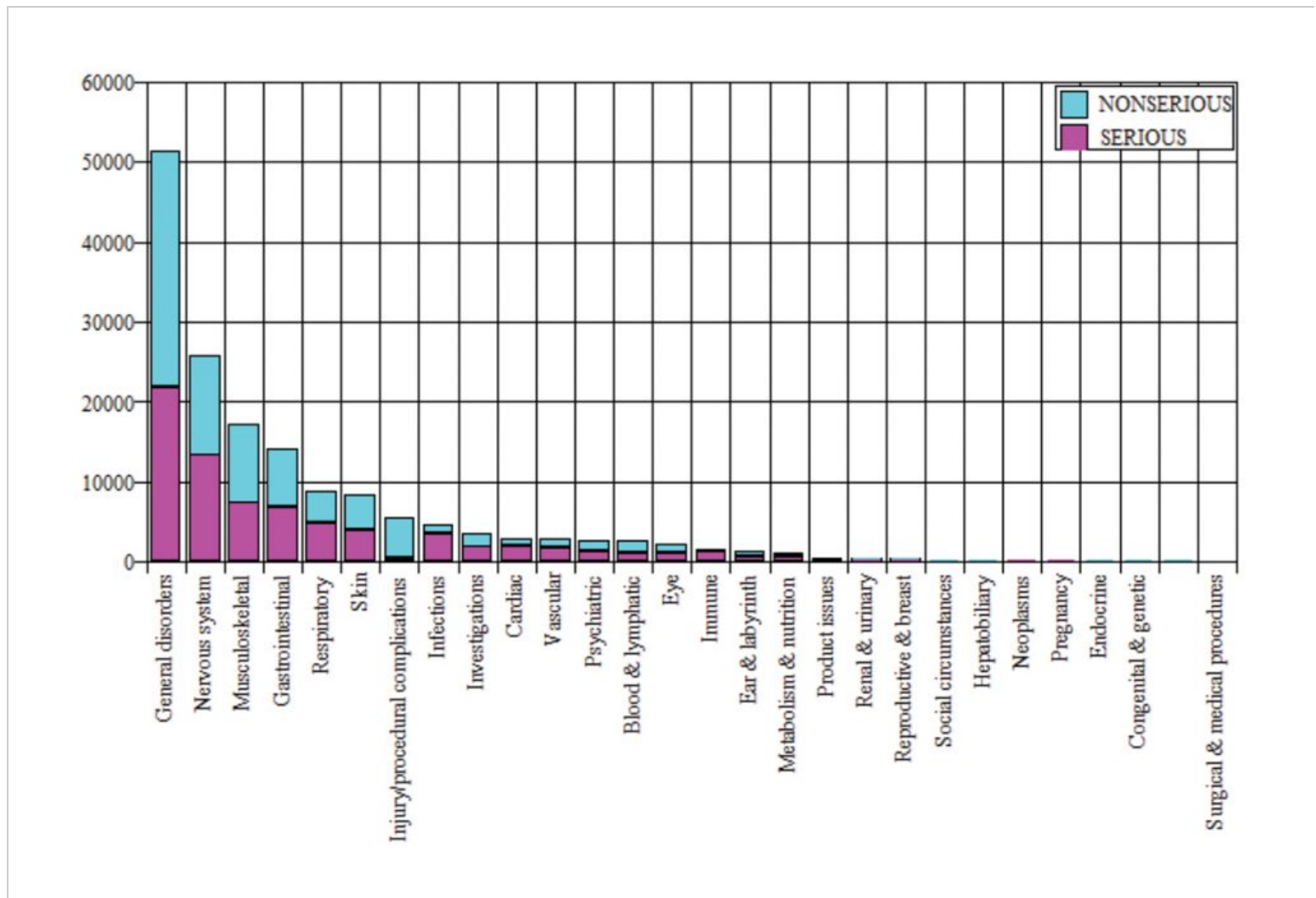


Figure 1: Total Numbers of BNT162b2 AEs by system organ classes and event seriousness

An overview of the characteristics of the recipients is given here:

Table 1. General Overview: Selected Characteristics of All Cases Received During the Reporting Interval

Characteristics		Relevant cases (N=42086)
Gender:	Female	29914
	Male	9182
	No Data	2990
Age range (years): 0.01 -107 years Mean = 50.9 years n = 34952	≤ 17	175 ^a
	18-30	4953
	31-50	13886
	51-64	7884
	65-74	3098
	≥ 75	5214
	Unknown	6876
Case outcome:	Recovered/Recovering	19582
	Recovered with sequelae	520
	Not recovered at the time of report	11361
	Fatal	1223
	Unknown	9400

a. in 46 cases reported age was <16-year-old and in 34 cases <12-year-old.

Of note, 1,223 recipients of the vaccine had a fatal outcome. More than 11,000 had not recovered. The outcome of 9,400 was unknown. Nearly three-quarters were female.

These numbers are concerning, but do they represent a significant safety concern? The answer to that question depends entirely upon the number of people who had been vaccinated up to that point.

Pfizer provided this number to the FDA in the general overview section of the document, section 3.1.1. — but in the document released under the FOIA request, that number was redacted:

“It is estimated that approximately (b) (4) doses of BNT162b2 were shipped worldwide from the receipt of the first temporary authorisation for emergency supply on 01 December 2020 through 28 February 2021.”

In the above, “(b)(4)” indicates that this number has been redacted.

The cumulative number of doses distributed worldwide as of Feb. 28 is not proprietary information, nor does it constitute personal, private data of individuals.

Yet without this key number there is no way to calculate the incidence of serious events, i.e., a safety signal.

The FDA chose, without explanation or any legal justification, to withhold this crucial piece of

data.

Despite the FDA's obvious intention to obfuscate, Pfizer provided a means of estimating this number when it unequivocally concluded: "... these signal detection analyses are consistent with the known safety profile of the vaccine."

What was the known safety profile of the vaccine?

As of Feb. 28, the only known safety profile of the vaccine was determined by the initial results from the [phase 3 trials](#) from the autumn of 2020.

Of 21,621 Pfizer vaccine recipients, 126 [[Polack FP, Thomas SJ, Kitchin N, et al., NEJM, Table S3](#)] suffered a serious adverse event in the trials. This is roughly one severe adverse event in 171.6 recipients.

Thus, if these data are consistent with its known safety profile, and roughly 79,000 serious adverse events had occurred up to that time, we can estimate that approximately 13,550,000 ($79,000 \times 171.6$) doses had been distributed.

Admittedly there is uncertainty in this calculation. Perhaps a different interpretation of the safety profile was implied.

However, Pfizer reported the number of doses that had been distributed, not administered.

Fewer doses would have been administered than delivered. Moreover, serious adverse events in the trials were distributed across participants who were fully vaccinated (having received two doses).

Urgent: 3 Ways to Help Stop Biden's Vaccine Mandates

Here we are using the number of doses as the denominator. This estimation will result in the lower limit of the true incidence of adverse events.

In other words, by using these assumptions we are giving Pfizer's vaccine the maximum benefit of the doubt.

Using this estimate of total doses given, the incidence of a fatal outcome is $1223/13.55$ million or 1 in 11,079.

Permanent sequelae (conditions that result as a consequence of vaccination) = $520/13.55$ million, or 1 in 26,057. Furthermore, 11,361 out of 13.55 million, or 1 in 1,193, had not yet recovered from an adverse event.

Pfizer inexplicably chose to group recipients who "recovered" with those who were "recovering."

How many in this large group (19,582) were still suffering from harm at the time of the report? On what basis did Pfizer determine a recipient still had a chance of full recovery?

With no clarification from the vaccine manufacturer, we are forced to lump them in with another large group of 9,400 whose ultimate outcome was “unknown” — leaving us with a high limit of 1 in 466 recipients having had an undetermined outcome.

Although none of these adverse events and fatalities were shown to be directly or indirectly caused by vaccination, Pfizer offered more data of concern around adverse events of “special Interest” (AESI).

According to Pfizer, 1,403 cardiovascular AESIs, 932 hematologic, 3,600 musculoskeletal, 501 neurologic and 3,674 “other” serious AESIs all occurred with a median time of onset of 24 hours or less from vaccination.

The 275 strokes and 449 cases of facial paralysis reported occurred with a median time of onset of two days from vaccination.

Though it is impossible to establish an unassailable causative link between vaccination and injury at this time, the temporal relationship between them is correlative and highly suggestive of causation.

Nevertheless, the authors of the Pfizer report concluded at the end of each AESI category that “This cumulative case review does not raise new safety issues.”

The report also included 24 serious cases in children younger than 12. Of those, 13 cases had not yet been resolved at the time of reporting. The mean age of these recipients was 3.7 years.

We must assume that very few children of that age were inoculated at that time given that Pfizer had authorization for use on adults only. With no number of inoculated children reported, we cannot know what the risk of injury is in children under 12.

Conclusions

Pfizer’s repeated assurances that no new safety issues exist are disingenuous at best.

The FDA was overtly obstructive by withholding crucial information required to make an accurate assessment of harm. However, by using reasonable estimations based on Pfizer’s own claims and published trial data, it is likely a safety signal does exist — and that safety signal was ignored by the very organization that is supposed to be listening for it, the FDA.

Pfizer’s estimated incidence of potential vaccine fatality, 1 in 11,079, is approximately twice that reported in VAERS. Given that the potential vaccine fatalities in this document have been passively reported, we can assume the actual incidence is higher.

More comprehensive [analyses](#) have demonstrated a VAERS underreporting factor of vaccine fatality approaching 41 or greater.

Underreported or not, the real and growing tragedy is that until an injury associated with vaccination is proven to be caused by it, it remains, for all intents and purposes, a non-existent signal to the very institutions responsible for public health and safety.

On what grounds can we as physicians and healthcare providers assure our patients this vaccine is safe if adverse events are not investigated or even acknowledged?

Is a nod from the FDA really good enough?

Or should we demand transparency, discussion or at the very least, unredacted data? What does the public expect of us?

SUGGEST A CORRECTION



Madhava Setty, M.D.

Madhava Setty, M.D. is senior science editor for The Defender.

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Name

**Ektor** • 3 months ago

Nothing to see here. Big pharma's motto is even if our poison mRNA injections murder millions, nothing will deter us from making \$100's of billions in profits. Nothing! There is no one to stop the carnage, death and misery.

23 ^ | v • Reply • Share ›

**jacques merde** → Ektor • 3 days ago

Please check out this document from D.A.R.P.A. just released by Project Veritas Look on Page 4 and you will see that the cure for COVID was known since early 2020 to be Ivermectin and HCQ.

Look on page one and it appears the USA (Fauci & Daszak) knew that SARS Cov 2 virus was created by Eco-Health Alliance in Wuhan, China.

JAG_Docs_pt1_Og_WATERMARK_OVER_Redacted.pdf (ctfassets.net)

Look at this document from the National Library of Medicine from 2005 after the first SARS Cov Virus outbreak.

They knew Chloroquine was the go-to drug for treatment.

Chloroquine is a potent inhibitor of SARS coronavirus infection and spread - PubMed (nih.gov)

D.A.R.P.A. when approached by Peter Daszak (Eco-Health Alliance) refused because the research was dangerously close to or was Gain Of Function which was not allowed in the States.

see more

3 ^ | v • Reply • Share ›

**Peter van der Stam** → Ektor • 2 days ago

And having seen a few videos of sports people crashing on the field and dying I have the next question:

Has anybody picked up that they are mainly black people dying on the field. Have they gotten a different contents in their jab??? There are different batch numbers available. One for whites and one for colored people?? I wonder

1 ^ | v • Reply • Share ›

**Sabrina Balsky** • 3 months ago

this is incomprehensible. Where is the regulatory agencies how is this even legal?

19 ^ | v • Reply • Share ›



CastleConnell → Sabrina Balsky • 3 months ago

The DOJ is captured, if it wasn't this would not be happening!

6 ^ | v • Reply • Share ›



Jacek R → CastleConnell • 4 days ago

Are they captured? Or are they bought out?

1 ^ | v • Reply • Share ›



Peter van der Stam → Jacek R • 2 days ago

BOTH !!

1 ^ | v • Reply • Share ›



Jan • 3 months ago

First, CDC and FDA defied public trust by working hand-in-hand with Pfizer to authorize the vaccine for emergency use and protect Pfizer from any injury liability; then last week, federal court approved Biden's administration's mandate to force it on almost every healthy person they manage to identify. The entire system has become invincible and indomitable.

17 ^ | v • Reply • Share ›



Cam → Jan • 3 months ago

Things aren't as we see them on the surface. the US Govt drove the agenda from the late 80's (both mRNA vaxes through DARPA, who called them, "population manipulation and species extinction technologies". COVID is a scam to cover up the vax and the vax is a weapon to put down the global population especially in those countries with large amounts of suppressed infectious diseases, such as Africa, Asia, India etc. the corporations (banking/central banksters, Big tech, Big Pharma, Big oil etc) have been pulling the strings of ever and they want more power just as the US Government wants to be head of the next NWO which will be a One world Government and One world Military, we are going to see a financial collapse to pave the way and probably a War with China to cement the deals. Just as we had WW1 then the league of nations WW2 and the UN, we will see a 3rd war and a new NWO. it's more about control then corruption and profits we are at war or at least under attack at the moment.

7 ^ | v • Reply • Share ›



Mick from Hooe → Jan • 3 months ago

FDA are allowing Pfizer 75 years to release ALL their dirty deadly secrets. The FDA only required 108 days to read the complete document themselves? This means they didn't even read the documents properly or they're in cahoots with Pfizer's desperation to stop

the public knowing the results. MAKE 'LIABILITY' the PRIMARY Condition of Full Approval and the Covid myth will disappear..

8 ^ | v 1 • Reply • Share ›



jacques merde → Mick from Hooe • 3 days ago

no, that is the CDC that wanted all that time and a judge said, "hell no!" here are the first batch of Pfizer docs...

<https://phmpt.org/pfizers-d...>

Enjoy!

2 ^ | v • Reply • Share ›



SEAN THOMAS • 3 months ago

"until an injury associated with vaccination is proven to be caused by it"

What would this require? Why don't we hold the "virus" up to the same standard?

14 ^ | v • Reply • Share ›



Mick from Hooe → SEAN THOMAS • 3 months ago

Any judge or adjudicator of DEAHs being caused directly by Vax will (disappear) or be silenced and replaced. That's today's world. FDA are allowing Pfizer 75 years to release ALL their dirty deadly secrets. The FDA only required 108 days to read the complete document themselves? This means they didn't even read the documents properly or they're in cahoots with Pfizer's desperation to stop the public knowing the results. MAKE 'LIABILITY' the PRIMARY Condition of Full Approval and the Covid myth will disappear.

3 ^ | v • Reply • Share ›



W. → Mick from Hooe • 3 months ago

More and more people in important public, political, professional positions are becoming aware of the reality that good effective medicines are being suppressed in order to legitimise this so-called vaccination. And unfortunately, they have so many bad side effects and brings deaths. Those in positions of responsibility do not want to be involved in this because they can no longer agree upon with their conscience and they are opting out or dropping out. Watch the video of the German Corona Committee, in which Dr. Reiner Füllmich gives a summary of the analyses of the last 1.5 to 2 years. It takes about 40 minutes, which is worthwhile. Send this email to all your friends, acquaintances, relatives, colleagues, anonymously or openly. Most people are good people and they have a conscience that guides them. And they believe in God, to whom they give many different names. And I believe in the good that will triumph. May the Primordial Consciousness help us.

3 ^ | v • Reply • Share ›

**Isthisthington** → W. • 3 months ago

Link ?

^ | v • Reply • Share ›

**jacques merde** → SEAN THOMAS • 3 days ago

shhhhh... (we know)

^ | v • Reply • Share ›

**shorebird** • 3 months ago

Until US hospitals fill with fully vaxxed patients due to ADE (or paradoxical immune system overreaction from exposure to a different or wild virus), no one is paying attention. After years of attempts, none, zero have resulted in a successful corona vaccn development, all run a serious risk of creating ADE, these "new" gene therapy shots are no exception. THE US FDA, Dr Fauci and others at the CDC and NIH are well aware of the history, yet denial is in abundance. Wait a month or two.

6 ^ | v • Reply • Share ›

**1555** → shorebird • 3 months ago

People are however paying attention to the MANY cruise ships, floating around with fully vaccinated crews, and passengers, yet FULL of Covid cases. How humorous that this is now being shown - for all to see.

2 ^ | v • Reply • Share ›

**jacques merde** → 1555 • 3 days ago

One cannot forget the Queen of England... Vaccinated 3 times but yet still caught COVID twice after vaccinations

1 ^ | v • Reply • Share ›

**Mick from Hooe** → shorebird • 3 months ago

Even when this inevitable day arrives the Vax makers will put out a new lie to deflect their responsibility. FDA are allowing Pfizer 75 years to release ALL their dirty deadly secrets. The FDA only required 108 days to read the complete document themselves? This means they didn't even read the documents properly or they're in cahoots with Pfizer's desperation to stop the public knowing the real intentions, contents and results. MAKE 'LIABILITY' the PRIMARY Condition of Full Approval and the Covid myth will disappear.

1 ^ | v • Reply • Share ›

**David Huber** • 3 months ago



I read the entire report two days ago. One of the things that struck me was their almost constant conclusion that "Conclusion: This cumulative case review does not raise new safety issues. Surveillance will continue" My take on this is that they knew that these things would happen and they were okay with these adverse events. So nothing to see, we [Pfizer] expected this. This is totally understandable as the Seneff & Nigh review of the "vaccine" research also stated that these events were possible.

Was there then informed consent? No. I would postulate that this then eliminates their indemnity from liability and should open them up for damages on a grand scale. They knew that their so called vaccine was dangerous, and hid that fact from the public. I also feel that there should be criminal charges brought against Pfizer, the FDA and the CDC.

One online group has been calling for an Office of the Inspector General of Health and Human Services investigation into the relationships and collusion, asking people to contact oig.hhs.gov to ask for this investigation. Since this call that part of the HHS web site has been down. This has been going on for at least 3 days. No oig.hhs.gov website available. Everything else on the HHS web site seems to be working, why this one part is not working and for so long seems rather ominous.

I will keep trying and I will also be petitioning my congressional representatives to take some corrective action.

5 ^ | v • Reply • Share ›



Mick from Hooe → David Huber • 3 months ago • edited

My (UK) MP has removed me from his emails - Huw Merriman hides from the truth and learns to say 'YES' to his lunatic controllers.

3 ^ | v • Reply • Share ›



eyesandears → David Huber • 3 months ago

David, Bingo! You hit the jackpot with your critical reading skills, which must be judiciously applied in the read or listen of everything. The defining word in the quote is "new" - heart, clot and bleed issues were fully known prior to forced market entry (traced years back thru patents and research from gov's own sites), making the incoming data anything but new.

As for criticality, my friend had Tucker on; Tuck made a statement which gave me a start concerning its meaning (I don't recall it accurately now). Said statement was then posted in writing, which I read and pointed out to my friend that Tuck had verbally conveyed the opposite of his ongoing publicized position, to even oppose all surrounding statements made in that episode. My friend stated, 'he really means this.' What's known is that in hearing a statement, one seeks info which confirms one's personal stance, and the brain will 'fill in' perceived gaps. A carefully worded statement may be heard to construe one view while its actuality conveys its opposing view. The twitter post on the above covered

view while in actuality espousing its opposing view. The written post on the screen served to concretely imprint in the mind of the reader the desired statement as a permanent register. Subtle but highly effective.

Charges? Who of those personally invested in said corps will work to destroy the income and gains derived? And the few who could do so may too often recall the senator who was beaten and hospitalized for non-melding views...

1 ^ | v • Reply • Share ›



Mick from Hooe → David Huber • 3 months ago

'informed consent' does nit exist in the UK either. it's an inconvenient opportunity for the public to learn the true potential of accepting a poison into their bodies.

1 ^ | v • Reply • Share ›



jacques merde → David Huber • 3 days ago

in the meanwhile, here is some reading material for ya! The firs rould of Pfizer docs released as court ordered...

<https://phmpt.org/pfizers-d...>

^ | v • Reply • Share ›



1555 → David Huber • 3 months ago

Pfizer even has a built in 'acceptable death' rate.

^ | v • Reply • Share ›



zark • 3 months ago

Look at the safety information they are recognizing now: Myocarditis and pericarditis occurs in some people who've received the vaccine, more commonly in males under 40 years of age than among females and older males.

So, this is saying that young men are getting heart inflammation. And what we know from all the anecdotal reports is 300 athletes have died or collapsed on the field, and children in schools have died of heart attacks.

And the reason they have to declare this is because they know it. They know it's happening. And the only way they can be sued is if they know there's a problem with their vaccine and they don't declare it. So, they declare it here, as if it's not that big of a deal.

2 ^ | v • Reply • Share ›



Formed • 3 months ago

This and the facebook articles are really well done and incredibly damning and significant. Well done guys

2 ^ | v • Reply • Share ›

**vinu arumugham** • 3 months ago

Financial/EMA reports show:

<https://www.sec.gov/Archive...>

More than 200 million doses of BNT162b2 supplied as of March 23, 2021

More than 450 million doses of BNT162b2 supplied to 91 countries or territories worldwide as of May 6, 2021

"The MAH estimated that approximately 415,922,715 doses of BNT162b2 were shipped worldwide from

the receipt of the first temporary authorisation for emergency supply on 1 December 2020 through 29

April 2021, corresponding to 332,738,172 estimated administered doses."

As of July 21, 2021, BioNTech and New York-based Pfizer, Inc. have shipped approximately one billion doses of BNT162b2 (Comirnaty) to more than 100 countries or territories worldwide.

May be useful to estimate the doses for Feb 28.

2 ^ | v • Reply • Share ›

**Mick from Hooe** → vinu arumugham • 3 months ago

Apparently there's a time limit coming up for 'Emergency Use' but that will be ignored by the Drug Barons and our Controllers.

2 ^ | v • Reply • Share ›

**Madhava Setty, MD** → vinu arumugham • a month ago

Thank you for sharing those numbers. Nevertheless, we are faced with the reality that knowing how many doses were shipped still won't tell us how many doses were injected. I believe the more concerning issue is that the FDA was willing to accept the number of doses shipped as the denominator. To me this demonstrates clear proof that they aren't really doing any oversight at all.

^ | v 2 • Reply • Share ›

**Ricky Ricardo** • 5 days ago

I have an internal document from Pfizer and I'm quite sure that I saw over 1,000 deaths in the trials. Regardless...they couldn't care less! When there is big money involved; human lives and suffering, isn't even a consideration!

1 ^ | v • Reply • Share ›

**jacques merde** → Ricky Ricardo • 3 days ago

here is the first batch of pfizer docs as court ordered... <https://phmpt.org/pfizers-d...>

2 ^ | v • Reply • Share ›



Kim from Brooklyn → jacques merde • 2 days ago

Thank you Jacques. I think.....I am going through them and my stomach is in knots. Did you see the video with Bourla (Pfizer) ? He is basically trying to blame the FDA, CDC, Fauci, etc. for "convincing" him to permit these jabs....now that the truth is out, isn't that what they do? Throw each other under the bus lickety split?

And where the heck is Fauci? I heard he is in a witness protection program.

^ | v • Reply • Share ›



Ricky Ricardo → jacques merde • 3 days ago

Thanks, Jacques. I'm just wondering if I will live long enough to even get through 10% of those documents!

^ | v • Reply • Share ›



jacques merde → Ricky Ricardo • 3 days ago

Ricky... If the court gets it's way... You will be suffocated in Pfizer documents... At least you will then know... Personally, I am royally disgusted by the actions of our governments, Big Pharma and the medical associations that lay down the rules that doctors have to follow... They all lost me when they denied "Natural immunity through infection"

1 ^ | v • Reply • Share ›



Ricky Ricardo → jacques merde • 2 days ago

That; not "the". I wish there was an edit button! Anyway; it's the first video on the page.

^ | v • Reply • Share ›



Ricky Ricardo → jacques merde • 2 days ago

Well; we won't have to go through 10%. In this CHD video; documents show the within one month of the trials, there were already over 1.000 deaths! I would call that significant - and of course, very evil!

<https://live.childrenshealt...>

^ | v • Reply • Share ›



Jan • 3 months ago

Universities are mandating booster shots due to omicron... but omicron is just a mild cold.

<https://www.stanforddaily.c...>

1 ^ | v • Reply • Share ›



jacques merde → Jan • 3 days ago

Jan here is some bathroom reading for you... the first batch of Pfizer docs as ordered from the courts...

<https://phmpt.org/pfizers-d...>

^ | v • Reply • Share ›



zark • 3 months ago

According to Dr Alix Mayer: By approving the Vaccine, the producers become liable for damages if they knew of the damages.

Vaccine makers love EUA emergency products because they have this huge liability shield. If you're injured by an EUA vaccine, you can't sue the manufacturer, you can't sue the person who gave it to you, you can't sue the institution where you got the shot.

Mandate for jabbing kids is their saver.

According to Mayer, the reason they're trying to mandate the COVID shot for children is to evade liability for injuries, because once a vaccine is on the childhood vaccination schedule, vaccine makers have immunity against lawsuits for injuries. Because then it gets full liability protection according to the 1986 Act. This is why they're going after our children when they have a 99.9973% recovery rate ...

1 ^ | v • Reply • Share ›



jacques merde → zark • 3 days ago

There is a huge difference in liability if the shot was given as "emergency use" or as a vaccine... If anyone is still getting covid shots, please ensure its the newer approved vaccine, not the "emergency use" supply... Drug companies are protected from liability using the "emergency use" jabs... Do your homework kids!

1 ^ | v • Reply • Share ›



Gary Brown • 3 months ago

December 20, 2021 School District in New York Sends out Email Warning Parents of Sudden Cardiac Arrest in Students Grades K-12

Eastport-South Manor Central School District (ESM) in Suffolk County, New York sent out an email to all parents informing them about the new physician in their district. The new hiring was part of the new regulations regarding Sudden Cardiac Arrest of students Grades K- 12.

<https://vaccineimpact.com/2...>

1 ^ | v • Reply • Share ›



CastleConnell → Gary Brown • 3 months ago

 Mind bending that it is acceptable to damage children routinely and viciously! What are the parents thinking?

3 ^ | v • Reply • Share ›



Gary Brown → CastleConnell • 3 months ago

They are not because of thier schooling or shall I say Indoctrination.

1 ^ | v • Reply • Share ›



jacques merde → CastleConnell • 3 days ago

sheep...



^ | v • Reply • Share ›



chris • 3 days ago

Pfizer 'only' produces their products, somebody else is 'approving' them for the general, completely misinformed or better yet, not informed at all public. Between those who approve and produce, whose fault is greater in the genetic massacre on human populations??? or is there a third invisible hand which guides it all?

^ | v • Reply • Share ›



Double_Up • 3 days ago

These stats are like saying only 2000 people have been killed in gun violence since 2000 in the U.S.

^ | v • Reply • Share ›

**anon** • 3 days ago

Watch this comprehensive religious/scientific/corrupt business video about history of vaccines, it has been removed from Youtube AND DuckDuckGo. These online modifiers mean business!

<https://odysee.com/@Confere...>

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**cpwise2003** • 3 days ago

I will forever ask why was it when the realization of the vaccines ridiculously poor efficacy became plainly evident. In the UK this was so during the period after the first two synchronized shots and well before the third shot was available.

Notwithstanding the false government narrative and censoring of spoken, written and portrayed challenges, I am asking just what research is being done or to be done about why it was so easy to have fooled such huge swathes of our nations to actually taken the third shot? I talk of the state of mind, susceptibility of these masses before conversion and why even blatant, in the face lies in the promotion of vaccines.

I am that much concerned, that sadly I can confidently say that before the word "covid" even starts to fade from the fore of memory, another blast of propaganda either for a fourth shot of same or a 'new' (nouvelle) pathogen will conveniently hit the news and that ALL the thrice jabbed will queue yet again without question and at least half of the twice jabbed will be once again sucked in to that propaganda vortex. Not to mention that "covid" has set the stage for an onslaught of vaccines for EVERY disease or ailment will be underway. Medical research will be solely that path and no other.

Its people to somehow have to change their passive outlook to an inquisitive one. How do we get the mass, clearly the majority, when confronted with such narratives to ask WHY, WHAT PURPOSE, WHERE IS THE DATA? (Show the workings to the public AS THE RESEARCH PROGRESSES and that INCLUDES ALL TRIAL DATA as a constant flow as it happens. Health of nations data should legally NEVER EVER require a F.O.I. request!!!

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